

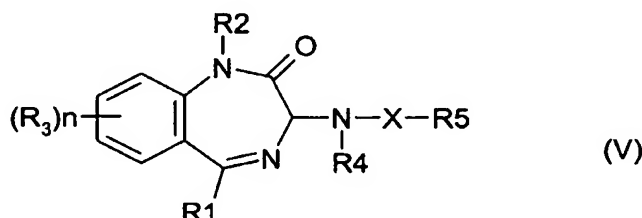
CLAIMS

1. A pharmaceutical composition which comprises a pharmaceutically acceptable carrier or diluent and:

- 5 (a) an inhibitor of the RSV fusion protein; and
(b) a benzodiazepine derivative capable of inhibiting RSV replication.

2. A composition according to claim 1, wherein component (b) is a compound of formula (V), or a pharmaceutically acceptable salt thereof,

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wherein:

- R^1 represents C_{1-6} alkyl, aryl or heteroaryl;
- 15 - R^2 represents hydrogen or C_{1-6} alkyl;
- each R^3 is the same or different and represents halogen, hydroxy, C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} alkylthio, C_{1-6} haloalkyl, C_{1-6} haloalkoxy, amino, mono(C_{1-6} alkyl)amino, di(C_{1-6} alkyl)amino, nitro, cyano, $-CO_2R'$, $-CONR'R''$, $-NH-CO-R'$, $-S(O)R'$, $-S(O)_2R'$, $-NH-S(O)_2R'$, $-S(O)NR'R''$ or $-S(O)_2NR'R''$, wherein each R' and R'' is the
- 20 same or different and represents hydrogen or C_{1-6} alkyl;
- n is from 0 to 3;
- R^4 represents hydrogen or C_{1-6} alkyl;
- X represents $-CO-$, $-CO-NR'-$, $-S(O)-$ or $-S(O)_2-$, wherein R' is hydrogen or a C_1-C_6 alkyl group; and
- 25 - R^5 represents an aryl, heteroaryl or heterocyclyl group which is substituted by a C_1-C_6 hydroxyalkyl group or a $-(C_1-C_4 \text{ alkyl})-X_1-(C_1-C_4 \text{ alkyl})-X_2-(C_1-C_4 \text{ alkyl})$ group, wherein

X₁ represents -O-, -S- or -NR', wherein R' represents H or a C₁-C₄ alkyl group and X₂ represents -CO-, -SO- or -SO₂-, or R₅ represents -A₁-Y-A₂, wherein:

- A₁ is an aryl, heteroaryl, carbocyclyl or heterocyclyl group;
- Y represents a direct bond or a C₁-C₄ alkylene, -SO₂-, -CO-, -O-, -S- or -NR' moiety, wherein R' is a C₁-C₆ alkyl group; and
- A₂ is an aryl, heteroaryl, carbocyclyl or heterocyclyl group.

3. A composition according to claim 2 wherein wherein R¹ is C₁₋₂ alkyl or phenyl.
- 10 4. A composition according to either claim 2 or claim 3, wherein wherein R² is hydrogen

5. A composition according to any one of claims 2 to 4 wherein R³ is halogen, hydroxy, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₁₋₄ alkylthio, C₁₋₄ haloalkyl, C₁₋₄ haloalkoxy, amino, 15 mono(C₁₋₄ alkyl)amino or di(C₁₋₄ alkyl)amino.

6. A composition according to claim 5 wherein R³ is fluorine, chlorine, bromine, C₁₋₂ alkyl, C₁₋₂ alkoxy, C₁₋₂ alkylthio, C₁₋₂ haloalkyl, C₁₋₂ haloalkoxy, amino, mono(C₁₋₂ alkyl)amino or di (C₁₋₂ alkyl)amino.

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7. A composition according to any of claims 2-6, wherein R⁴ is hydrogen or C₁₋₂ alkyl.

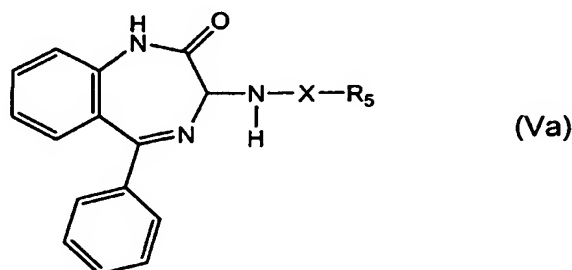
8. A composition according to any one of claims 2-7, wherein X is -CO- or -CO-NR' wherein R' represents hydrogen or a C₁-C₂ alkyl group.

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9. A composition according to any one of claims 2-8, wherein R⁵ is a 5- or 6- membered heterocyclyl, aryl or heteroaryl ring which is substituted by a C₁-C₆ hydroxyalkyl group or a -(C₁-C₄ alkyl)-X₁-(C₁-C₄ alkyl)-X₂-(C₁-C₄ alkyl) group, wherein X₁ and X₂ are as defined in 30 claim 2.

10. A composition according to claim 9, wherein R⁵ is a 5- or 6- membered heteroaryl group which is substituted by a -CH₂-OH or -(C₁-C₄ alkyl)-NR'-(C₁-C₄ alkyl)-S(O)₂-(C₁-C₄ alkyl) substituent, wherein R' is hydrogen or C₁-C₂ alkyl.
- 5 11. A composition according to claims 2-10, wherein A₁ is an aryl or heteroaryl group.
12. A composition according to claim 11, wherein A₁ is a phenyl group, a monocyclic 5- or 6- membered heteroaryl group or a 5- to 6- membered heteroaryl group fused to a monocyclic oxo-substituted 5- to 6- membered heterocyclyl group.
- 10 13. A composition according to claims 2-12 wherein A₁ is unsubstituted or substituted by 1 or 2 substituents selected from halogen, cyano, nitro, C₁-C₄ alkyl, C₁-C₄ haloalkyl and C₁-C₄ alkoxy substituents.
- 15 14. A composition according to claims 2-13, wherein Y represents a direct bond, a C₁-C₂ alkylene group, -SO₂- or -O-.
15. A composition according to claims 2-14 wherein A₂ is a phenyl, 5- to 6- membered heteroaryl, 5- to 6- membered heterocyclyl or C₃-C₆ cycloalkyl group.
- 20 16. A composition according to claims 2-15, wherein when A₂ is a heterocyclyl group it is attached to the moiety Y via a N atom.
17. A composition according to claims 2-16, wherein A₂ is unsubstituted or is substituted
- 25 by 1 or 2 substituents which are selected from C₁-C₄ alkyl and halogen substituents when A₂ is a heteroaryl or aryl group and which are selected from C₁-C₄ alkyl, halogen and oxo substituents when A₂ is a carbocyclic or heterocyclyl group.
18. A composition according to claims 2-17, wherein A₂ is a piperazinyl, pyridyl,
- 30 morpholinyl, pyrrolidinyl, piperidinyl, pyrazinyl, cyclopropyl, phenyl or S,S-dioxo-thiomorpholino group, which is unsubstituted or substituted by a C₁-C₂ alkyl group.

19. A composition according to any one of claims 2-18 wherein the benzodiazepine derivative of formula (V) is a benzodiazepine derivative of formula (Va):



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wherein:

- X is -CO- or -CO-NH-; and
- R⁵ is a 5- to 6- membered heteroaryl group, for example a furanyl group,
 10 which is substituted by -CH₂-OH or -(C₁-C₄ alkyl)-N(CH₃)-(C₁-C₄ alkyl)-SO₂-(C₁-C₄ alkyl)
 or R₅ represents -A₁-Y-A₂, wherein:
 - A₁ is a phenyl, pyridyl, furanyl, thiazolyl, oxazolyl, isoxazolyl, thienyl or
 1H-imidazo[4,5-b]pyridin-2-(3H)-one moiety, which is unsubstituted or substituted by 1 or 2
 substituents selected from halogen, cyano, C₁-C₂ alkyl, C₁-C₂ haloalkyl and C₁-C₂ alkoxy
 15 substituents;
 - Y is a direct bond, a C₁-C₂ alkylene group, -SO₂- or -O-; and
 - A₂ is a piperazinyl, pyridyl, morpholinyl, pyrrolidinyl, piperidinyl, pyrazinyl,
 cyclopropyl, phenyl or S,S-dioxo-thiomorpholino group, which is unsubstituted or
 substituted by a C₁-C₂ alkyl group.

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20. A composition according to claim 1, wherein the benzodiazepine derivative of formula (V) is:

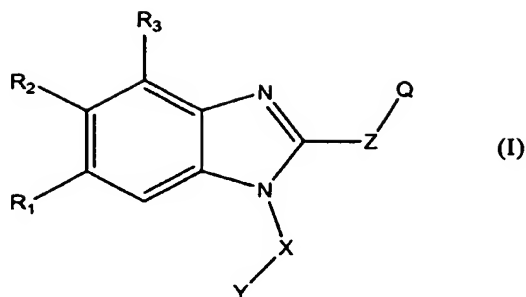
- 6-(4-Methyl-piperazin-1-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]
 diazepin-3-yl)-nicotinamide;
- 25 3,4,5,6-Tetrahydro-2H-[1,2']bipyridinyl-5'-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-
 benzo[e][1,4]diazepin-3-yl)-amide;

- (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-2-Chloro-4-morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- 5 (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-4-fluoro-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-5-Chloro-2-(1,1-dioxo-1 λ 6-thiomorpholin-4-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-5-fluoro-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- 10 (S)-5-(4-Methyl-piperazin-1-ylmethyl)-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-5-Pyrrolidin-1-ylmethyl-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 15 (S)-5-Piperidin-1-ylmethyl-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-5-Dimethylaminomethyl-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-4-Fluoro-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-2-piperidin-1-yl-benzamide;
- 20 (S)-4-Fluoro-2-morpholino-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-4-Cyano-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-2-pyrrolidin-1-yl-benzamide;
- 25 (S)-4-Cyano-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-piperidine-1-yl-benzamide;
- (S)-N-(2-Oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-2-pyrrolidin-1-yl-4-trifluoromethyl-benzamide;
- (S)-N-(2-Oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-2-piperidin-1-yl-4-trifluoromethyl-benzamide;
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- (S)-2-Morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-4-trifluoromethyl-benzamide;
- (S)-N-(2-Oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-2-pyrrolidin-1-yl-5-trifluoromethyl-benzamide;
- 5 (S)-2-Morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-5-trifluoromethyl-benzamide;
- (S)-2-Morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-nicotinamide;
- (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-
- 10 benzo[e][1,4]diazepin-3-yl)-nicotinamide;
- (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-2-methyl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-4-methyl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- 15 (S)-2-(1,1-Dioxo-1 λ 6-thiomorpholin-4-yl)-6-methyl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-2-Chloro-6-(1,1-dioxo-1 λ 6-thiomorpholin-4-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-3-Cyclopropyl-2-oxo-2,3-dihydro-imidazo[4,5-b]pyridine-1-carboxylic acid (2-oxo-5-
- 20 phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-3-(4-Methyl-piperazine-1-sulfonyl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-4-(4-Methyl-piperazin-1-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- 25 (S)-N-(2-Oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-3-(piperidine-1-sulfonyl)-benzamide;
- (S)-3-(Morpholine-4-sulfonyl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-5-Morpholin-4-ylmethyl-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-
- 30 benzo[e][1,4]diazepin-3-yl)-amide;

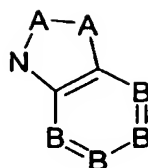
- (S)-5-Hydroxymethyl-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-5-(1,1-Dioxo-1 λ 6-thiomorpholin-4-ylmethyl)-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 5 (S)-2-Chloro-4-(1,1-dioxo-1 λ 6-thiomorpholin-4-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-2-Chloro-5-(1,1-dioxo-1 λ 6-thiomorpholin-4-yl)-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-5-{[(2-Methanesulfonyl-ethyl)-methyl-amino]-methyl}-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 10 (S)-2-Pyridin-3-yl-thiazole-4-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-2-Pyridin-4-yl-thiazole-4-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 15 (S)-4-Methyl-2-pyrazin-2-yl-thiazole-5-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-2-Morpholin-4-ylmethyl-furan-3-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-3-Morpholin-4-ylmethyl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- 20 (S)-5-Morpholin-4-ylmethyl-isoxazole-3-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-3-Morpholin-4-ylmethyl-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 25 (S)-5-Pyridin-2-yl-thiophene-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-2-Methyl-4-(morpholin-4-sulfonyl)-furan-3-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-6-Morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-nicotinamide;
- 30

- (S)-3-Morpholin-4-ylmethyl-thiophene-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- (S)-5-Morpholin-4-ylmethyl-thiophene-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 5 2-Morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide;
- (S)-5-Phenyl-oxazole-4 carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide;
- 1-(2-Oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-3-(4-phenoxy-phenyl)-urea
- 10 an N-oxide of any of the above compounds;
- or a pharmaceutically acceptable salt thereof.
21. A composition according to claim 1, wherein the benzodiazepine derivative of formula (V) is (S)-5-(1,1-Dioxo-1λ6-thiomorpholin-4-ylmethyl)-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide or (S)-2-Chloro-4-morpholin-4-yl-N-(2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-benzamide
- 15 or a pharmaceutically acceptable salt thereof.
22. A composition according to claim 21, wherein the benzodiazepine derivative of formula (V) is (S)-5-(1,1-Dioxo-1λ6-thiomorpholin-4-ylmethyl)-furan-2-carboxylic acid (2-oxo-5-phenyl-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl)-amide or a pharmaceutically acceptable salt thereof.
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23. A composition according to any one of the preceding claims wherein component
- 25 (a) is a compound of formula (I), or a pharmaceutically acceptable salt thereof,



wherein:

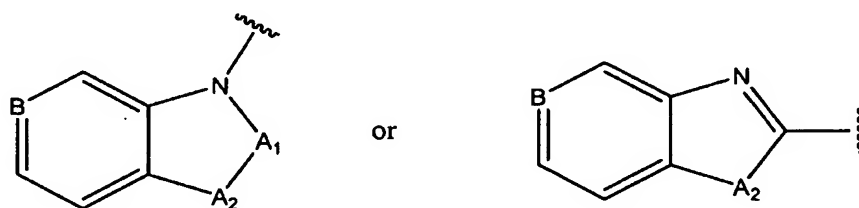
- 5 - X is H or C₁₋₆ alkyl; said C₁₋₆ alkyl being optionally substituted with halogen, OCOR₄ or S(O)_n-C₁₋₆ alkyl;
- Y is R₄, NR₄R₅, NCOR₄, =N-OR₄, -CONHR₄, COOR₄, -OR₄, aryl, heteroaryl, cyclyl or heterocyclyl, where R₄ and R₅ are H or C₁₋₆ alkyl;
- Z is CR₆R₇, where R₆ and R₇ are independently H, or straight, branched or cyclic C₁₋₆ alkyl;
- 10 - n is 1-6;
- R₁ is CONR₄R₅, CO₂R₄ or C₁₋₆ alkyl, said C₁₋₆ alkyl can be optionally substituted with OR₄ or NR₈R₉;
- R₈ and R₉ are each independently H, C₁₋₆ alkyl, SO₂R₅, CO₂R₄ or COR₄;
- 15 - R₂ is selected from the group consisting of NH₂, CONR₆R₇, heteroaryl, C₂₋₆ alkenyl, CO₂R₄, N=CPh₂, C(=NH)NH₂ and C₁₋₆ alkyl; said alkyl optionally substituted with a member selected from the group consisting of halogen, CN, NR₁₀R₁₁, OSO₂R₄ and OR₄;
- R₉ and R₁₀ are each independently selected from the group consisting of H, C₁₋₆ alkyl,
- 20 - C₃₋₆ cycloalkyl, CO₂R₄, COR₄ and SO₂R₄;
- R₃ is selected from the group consisting of (1) CO₂R₉; (2) C₁₋₆ alkyl optionally substituted with CN, OR₄ or NR₆R₇; and (3) C₂₋₆ alkenyl substituted with CN;
- Q is a member selected from the group consisting of



A is C or N, optionally substituted with H, halogen, straight, branched or cyclic C₁₋₆ alkyl, C₂₋₆ alkenyl, CO₂R₄, aryl or C₃₋₆ cycloalkyl. Where A is carbon, it may also be
 5 optionally substituted by O or S via a double bond;

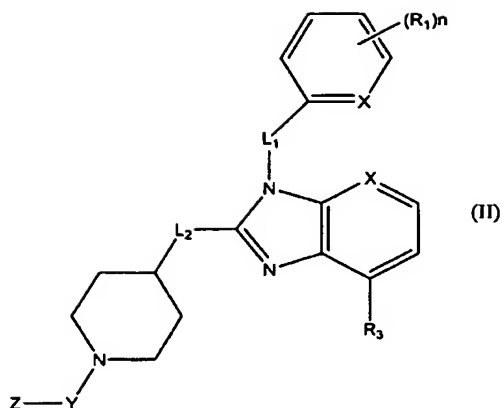
B is C or N; where B is C it may be optionally substituted by H, C₁₋₆ alkyl, NO₂, CN, halogen, COR₄, COOR₄, CONHR₄C(=NH)NH₂ or C(=NOH)NH₂.

24. A composition according to claim 23 wherein component (a) is a compound of
 10 general formula (I), as defined above, or a pharmaceutically acceptable salt thereof, wherein at least two of R₁, R₂ and R₃ are hydrogen, and the other is hydrogen or -C(NH)-NH₂ and/or -X-Y is H, or X is a C₁₋₆ alkylene group which is unsubstituted or substituted by a hydroxy group and Y is H, OH, CN, -NR'R'', -COR', -SO₂R' or phenyl, wherein R' and R'' are the same or different and represent a C₁₋₄ alkyl group and/or Z is -CH₂- and/or Q is a moiety
 15



wherein B is -CH- or -N-, A₁ is -C(O)- or -NH- and A₂ is -CH₂-, -CHR'- or -NR''-,
 wherein R' is a halogen atom and R'' represents a hydrogen atom or a C₁₋₄ alkyl, C₂₋₄
 20 alkenyl, C₃₋₆ cycloalkyl, -SO₂-(C₁₋₆ alkyl), -SO₂-N(C₁₋₆ alkyl)₂ or -(CO-NH)_a-(C₁₋₄ alkyl)-phenyl group, wherein a is 0 or 1, which group is unsubstituted or is substituted with a hydroxy or cyano substituent.

25. A composition according to claims 1 to 22 wherein component (a) is a compound of
 25 formula (II), or a pharmaceutically acceptable salt thereof,



wherein:

- 5 - L_1 is $-\text{CH}_2-$ or $-\text{CHR}_2-\text{CO}-$
- each X is the same or different and CH or N;
- each R_1 is the same or different and is C_{1-6} alkyl, halogen, hydroxy, phenyl or $(\text{CH}_2)_m=\text{NH}_2$;
- n is 1 or 2;
- 10 - R_2 is C_{1-6} alkoxy or C_{1-6} alkoxy-phenyl;
- R_3 is C_{1-6} alkyl;
- L_2 is $-\text{CH}_2-$ or $-\text{NH}-$;
- Y is C_{1-6} alkyl or C_{1-6} alkenyl;
- Z is H, $\text{N}(\text{R}_4)_2$, $-\text{C}(=\text{O})-\text{R}_5$, $-\text{C}(=\text{CH}_2)-\text{R}_5$, $-\text{CH}(\text{OH})-\text{R}_5$, $-\text{CH}(\text{CH}_3)-\text{R}_5$, $-\text{CH}(\text{OCH}_3)-$
- 15 R_5 ;
- each R_4 is the same or different and is H, C_{1-6} alkyl;
- R_5 is C_{1-6} alkyl-carbonyl, amino, hydroxyl, aryl, heteroaryl, carbocyclyl, heterocyclyl;
- and
- $m=1-6$

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26. A composition according to anyone of claims 1 to 22, wherein component (a) is:
 1-Cyclopropyl-3-[1-(4-hydroxy-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-imidazo[4,5-c]pyridin-2-one

- {2-[2-(1,2-Dihydro-benzotriazol-1-ylmethyl)-benzoimidazol-1-yl]]ethyl}-diethyl-amine
- {2-[2-(3-Iodo-2,3-dihydro-indazol-1-ylmethyl)-benzimidazol-1-yl]-ethyl}-dimethyl-amine
- 5 1-Isopropenyl-3-[1-(3-methyl-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-benzoimidazol-2-one
- 1-(4-Hydroxy-benzyl)-3-[1-(3-methyl-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-benzoimidazol-2-one
- 1-Isopropenyl-3-[1-(3-oxo-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-benzoimidazol-2-one
- 10 1-Ethyl-3-[1-(2-hydroxy-2-phenyl-ethyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-benzoimidazol-2-one
- 1-Ethyl-3-[1-(4-hydroxy-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-benzoimidazol-2-one
- 15 7-[2-(3-Isopropenyl-2-oxo-2,3-dihydrobenzoimidazol-1-ylmethyl)-benzoimidazol-1-yl]-heptanenitril
- 5-{3-[1-(3-Methanesulfonyl-propyl)-1H-benzoimidazol-2-ylmethyl]-2-oxo-2,3-dihydro-benzoimidazol-1-yl}-pentanenitrile
- 3-[1-(3-Methyl-butyl)-1H-benzoimidazol-2-ylmethyl]-2-oxo-2,3-dihydro-benzoimidazol-1-carboxylic acid benzylamide
- 20 1-Methanesulfonyl-3-[1-(3-methyl-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-benzoimidazol-2-one
- 3-[1-(3-Methyl-butyl)-1H-benzoimidazol-2-ylmethyl]-2-oxo-2,3-dihydro-benzoimidazol-1-sulfonic acid dimethylamide
- 25 1-Isopropenyl-3-(1-propyl-1H-benzoimidazol-2-ylmethyl)-1,3-dihydro-imidazo[4,5-c]pyridine-2-one
- Bis(5-amidino-2-benzimidazolyl)-methane
- 2-{2-[1-[1-(2-Amino-ethyl)-piperidin-4-ylamino]-4-methyl-benzoimidazol-1-ylmethyl]-6-methyl-pyridin-3-ol
- 30 or a pharmaceutically acceptable salt thereof.

27. A composition according to any one of claims 1 to 22, wherein component (a) is 1-cyclopropyl-3-[1-(4-hydroxy-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-imidazo[4,5-c]pyridin-2-one, {2-[2-(1,2-dihydro-benzotriazol-1-ylmethyl)-benzoimidazol-1-yl]ethyl}-diethyl-amine, {2-[2-(3-iodo-2,3-dihydro-indazol-1-ylmethyl)-benzimidazol-1-yl]-ethyl}-dimethyl-amine or a pharmaceutically acceptable salt thereof.

28. A composition according to any one of claims 1 to 22, wherein component (a) is 1-cyclopropyl-3-[1-(4-hydroxy-butyl)-1H-benzoimidazol-2-ylmethyl]-1,3-dihydro-imidazo[4,5-c]pyridin-2-one or 1-Isopropenyl-3-(1-propyl-1H-benzoimidazol-2-ylmethyl)-1,3-dihydro-imidazo[4,5-c]pyridine-2-one or a pharmaceutically acceptable salt thereof.

29. A composition according to any one of the preceding claims wherein component (a) is present in an amount of from 0.025 wt% to 10 wt%.

30. A composition according to any one of the preceding claims wherein component (b) is present in an amount of 0.025 wt% to 10 wt%.

31. A composition according to any one of the preceding claims, for use in the treatment of the human or animal body.

32. Use of:

(a) an RSV fusion protein inhibitor as defined in any one of claims 1 and 23 to 28; and

(b) a benzodiazepine derivative defined in any one of claims 1 to 22,

in the manufacture of a medicament for use in treating or preventing an RSV infection.

33. Use according to claim 32, wherein the medicament is a composition as defined in claim 29 or 30.

34. A product comprising:

- (a) an RSV fusion protein inhibitor as defined in any one of claims 1 and 23 to 28; and
- (b) a benzodiazepine derivative as defined in any one of claims 1 to 22;
- for separate, simultaneous or sequential use in the treatment of the human or animal body.
- 5
35. A product according to claim 34 for separate, simultaneous or sequential use in treating or preventing an RSV infection.
36. A method of treating or preventing an RSV infection in a patient, which method
- 10 comprises the administration to said patient of:
- (a) an RSV fusion protein inhibitor as defined in any one of claims 1 and 23 to 28; and
- (b) a benzodiazepine derivative as defined in any one of claims 1 to 22.
- 15 37. Use of an RSV fusion protein inhibitor as defined in any one of claims 1 and 23 to 28, in the manufacture of a medicament for use in treating or preventing an RSV infection, by co-administration with a benzodiazepine derivative as defined in any one of claims 1 to 22.
38. Use of a benzodiazepine derivative as defined in any one of claims 1 to 22, in the
- 20 manufacture of a medicament for use in treating or preventing an RSV infection, by co-administration with an RSV fusion protein inhibitor as defined in any one of claims 1 and 23 to 28.